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August 2, 2011



Via Federal Express

United States Environmental Protection Agency - East Attn: TSCA Section 8(e) Room 6428 1201 Constitution Avenue, NW Washington, DC 20004

Subject:

Notice in Accordance with Section 8(e): Results of a Test Study in Female, Non-

Pregnant Wistar Rats with [

Dear Sir/Madam:

] is submitting	results of a T	est Study in Fe	emale, Non-Pregn	ant Wistar Rats
[Crl:WI(HAN)] with [
], cond	ucted by []. The substance
is a developmental p	esticide.				

The test substance was administered to groups of 3 non-pregnant Wistar rats via gavage for 14 days. On day 14, all surviving animals were sacrificed and examined.

Following dose levels had been investigated:

50, 100, 200 and 400 mg/kg bw/d

The following findings were observed (in comparison to control):

400 mg/kg bw/d

- Poor general state starting after 2 days of exposure (increasing incidence or severity of reduced nutritional state, piloerection, lateral position, unsteady gait, and apathy)
- Reduced food consumption (down to 46%)
- Individual body weight loss (up to -9%)
- Sacrificed moribund (3/3 animals on day 5 of the study)

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United States Environmental Protection Agency - East August 2, 2011 Page 2

200 mg/kg bw/d

- Poor general state starting after 3 days of exposure (increasing incidence or severity of piloerection, hunched posture, and ataxia after treatment)
- Reduced food consumption (down to 49%)
- Individual body weight losses (up to -11%)
- Sacrificed moribund (1/3 animals on day 6 and 1/3 animals on day 12 of the study)

100 and 50 mg/kg bw/d

No treatment-related findings

[accordance with EPA's policy.

Please note that a confidential version of this letter is enclosed, treating the chemical identity and company identity as Confidential Business Information.

A Confidentiality Substantiation Questionnaire is being submitted.

Sincerely,

Enclosures